

I. AMENDMENTS

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

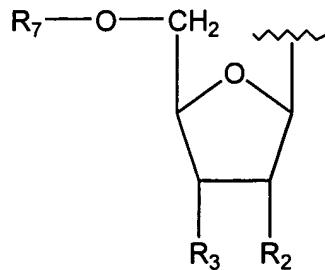
Claims 1 – 55. (Canceled).

56. (Currently Amended) A method for inhibiting the proliferation of a hyperproliferative neoplastic cell that endogenously overexpresses thymidylate synthase, comprising contacting the cell with a compound of claim 62 or a 5'-monophosphate metabolite thereof formed after administration to a subject.

57. (Currently Amended) A method for treating a pathology characterized by hyperproliferative neoplastic cells that endogenously overexpresses thymidylate synthase in a subject comprising administering to the subject a compound of claim 62 or a 5'-monophosphate metabolite thereof formed after administration to a subject.

58. (Canceled).

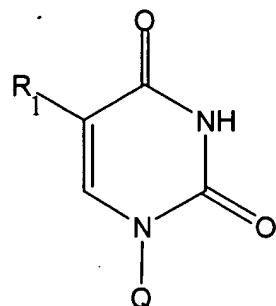
59. (Previously Presented) The method of claim 56 or 57, wherein Q has the formula:



60. (Previously Presented) The method of claim 56 or 57, wherein R_1 is a halogen.

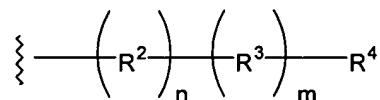
61. (Previously Presented) The method of claim 56 or 57, wherein R₁ is of the formula (-CH=CH)_n-R₄, wherein n is an integer from 1 to 10, and R₄ is selected from the group consisting of H; hydroxyl; a halogen; -NHCHO; -OCN; -SCN; -N₃; -NH₂; -NHOH; -NHNH₂ and a C₂ to C₄ carbon-containing substituent selected from the group consisting of alkyl, alkenyl, alkynyl, -O-alkyl, -O-aryl, O-heteroaryl, -S-alkyl, -S-aryl, -S-heteroaryl, -NH-alkyl, -N(alkyl)₂ and NHO-alkyl.

62. (Currently Amended) A compound of the formula:



wherein:

R¹ is of the formula:

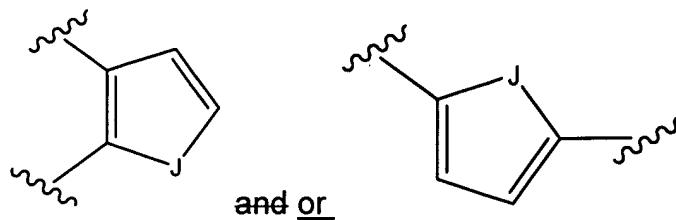


wherein R² is one of:

an unsaturated C₂ to C₄ hydrocarbyl group;

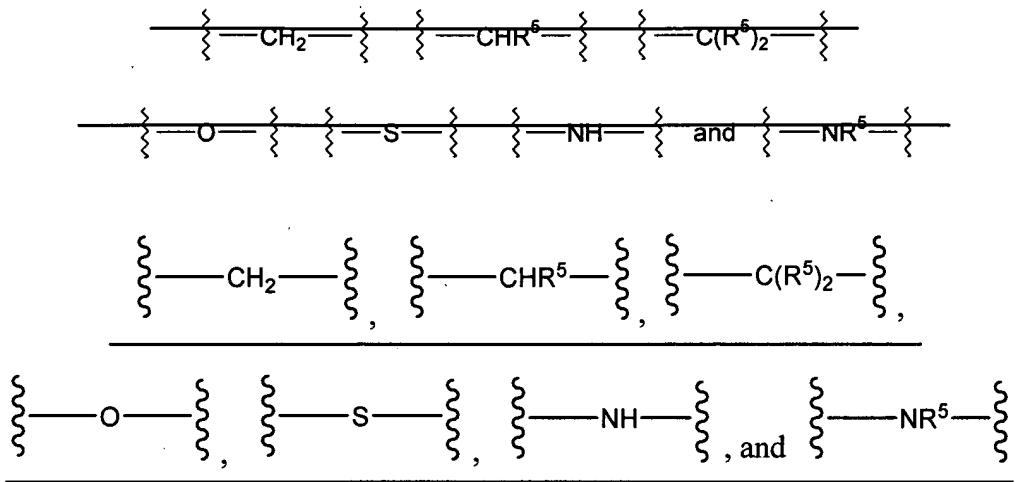
~~an aromatic C₄-X hydrocarbyl group, wherein X is the heteroatom;~~ or

a heteroaromatic group having the structure:



wherein J is -O-, -S-, -Se-, -NH-, or -NR^{ALK}-, wherein R^{ALK} is a linear or branched alkyl having 1 to 10 carbon atoms or a cycloalkyl group having 3 to 10 carbon atoms;

R³ is selected from the group consisting of:

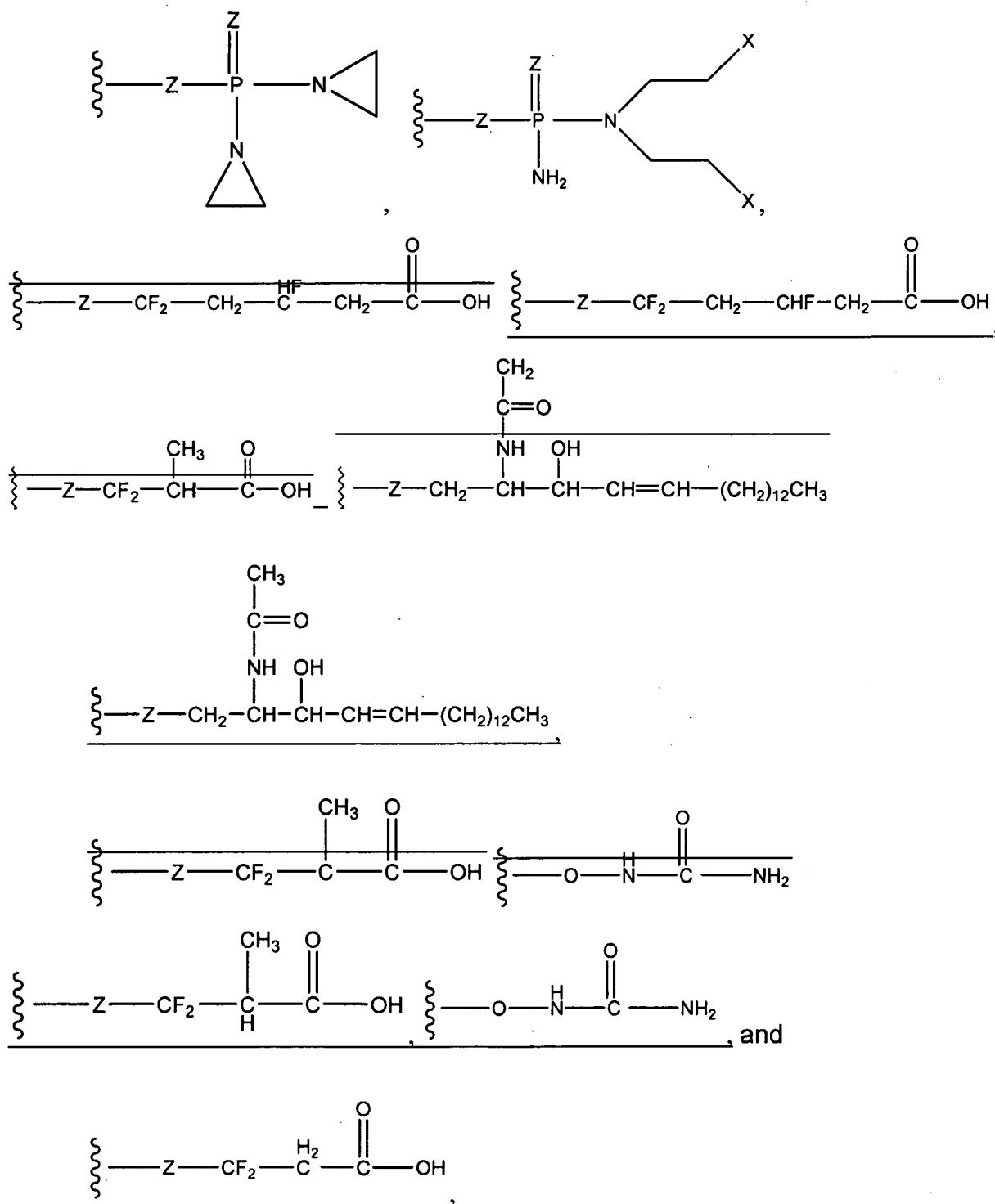


wherein R⁵ may be the same or different and is independently a linear or branched alkyl group having from 1 to 10 carbon atoms, or a cycloalkyl group having from 3 to 10 carbon atoms;

wherein n is an integer from 1 to 10;

wherein m is 0 or 1;

wherein R⁴ is a toxophore selected from the group consisting of:



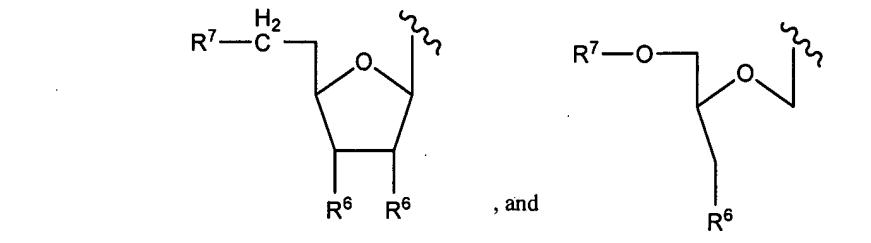
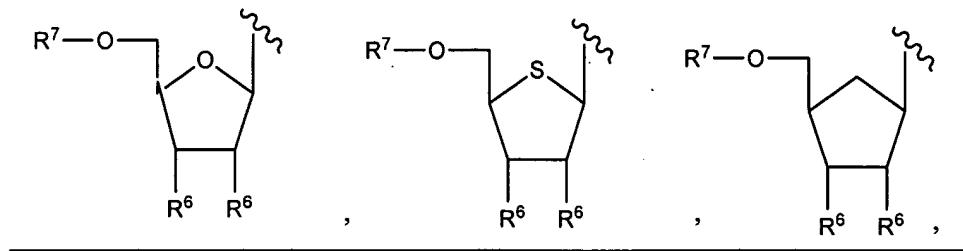
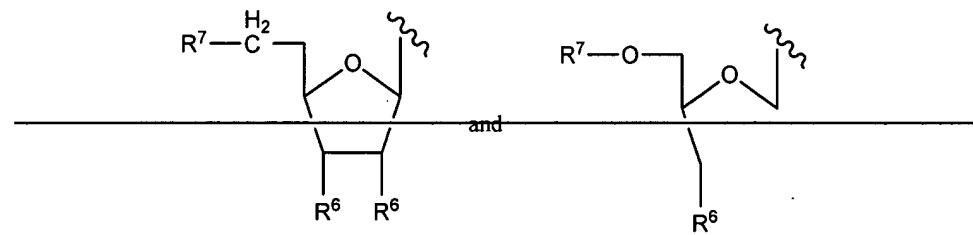
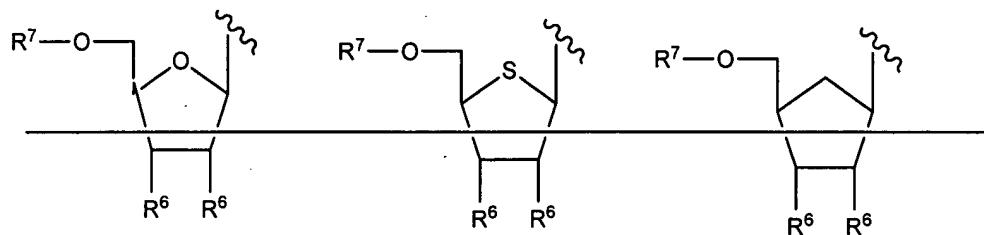
wherein X is -Cl, -Br, -I, or other halogen, with the proviso that when R⁷ is -H, and

~~M is zero m is zero~~, then R^4 is not a halogen or when m is zero and n is zero, then R^4 is not a halogen;

wherein Y is independently -H or -F;

wherein Z is independently -O- or -S-;

wherein Q is selected from the group consisting of:

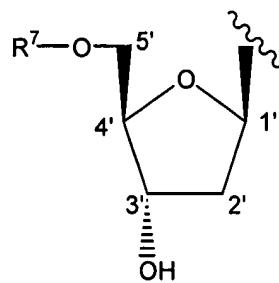


wherein R⁶ is independently -H, -OH, -OC(=O)CH₃, or -O-R₉ wherein R₉ is a hydroxyl protecting group other than acetyl; and,

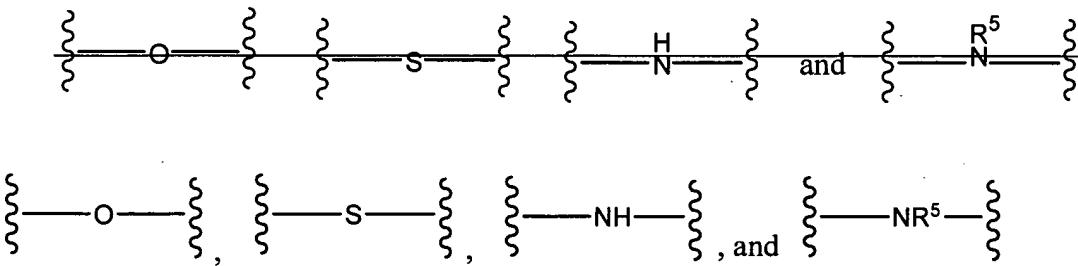
wherein R⁷ is hydrogen or a phosphoramidatyl derivative of a naturally-occurring amino acid;

and wherein said compound may be in any enantiomeric, diastereomeric, or stereoisomeric form, consisting of a D-form, L-form, α-anomeric form, and β-anomeric form.

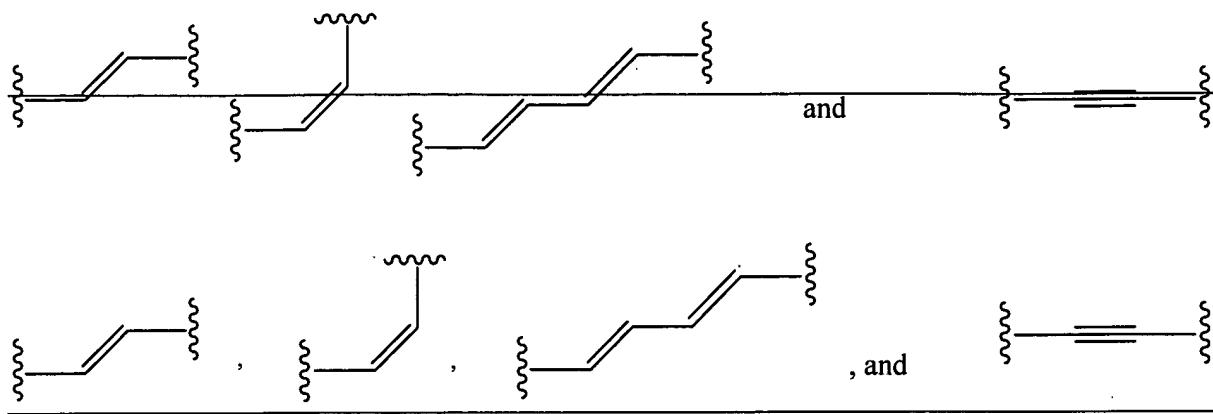
63. (Original) A compound according to claim 62, wherein Q is:



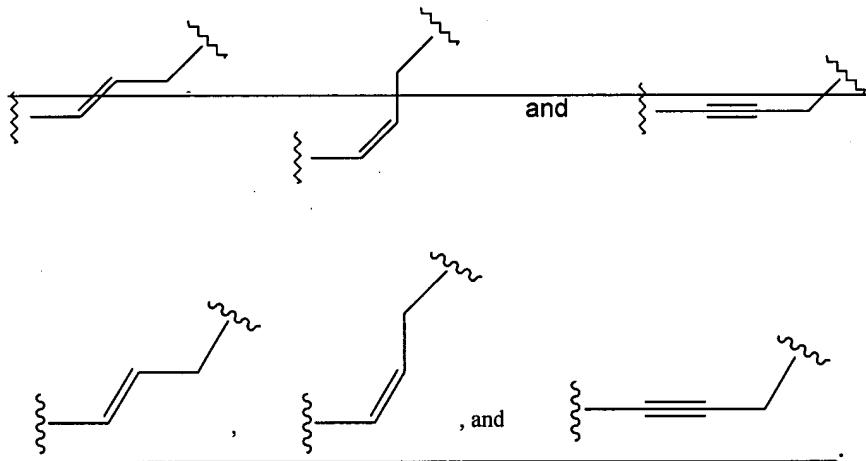
64. (Currently Amended) A compound of claim 62, wherein R³ is selected from the group consisting of:



65. (Currently Amended) A compound of claim 62, wherein R² is selected from the group consisting of:



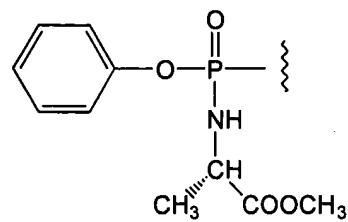
66. (Currently Amended) A compound of claim 62, wherein R^2 and R^3 , taken together form a structure selected from the group consisting of:



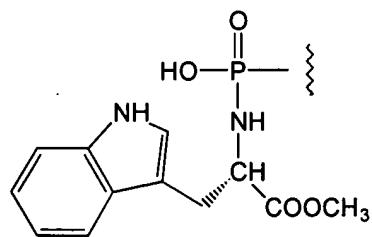
67. (Canceled).

68. (Canceled).

69. (Previously Presented) A compound of claim 62, wherein R^7 is:



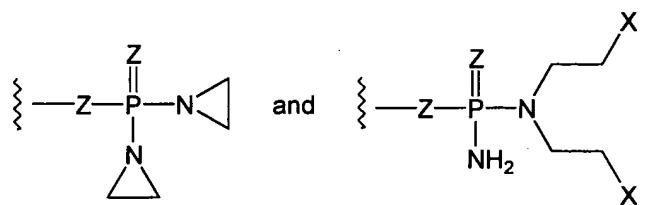
70. (Previously presented) A compound of claim 62, wherein R⁷ is:



71. (Canceled).

72. (Canceled).

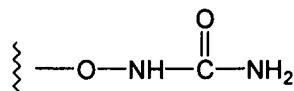
73. (Original) A compound of claim 62, wherein R⁴ is selected from the group consisting of:



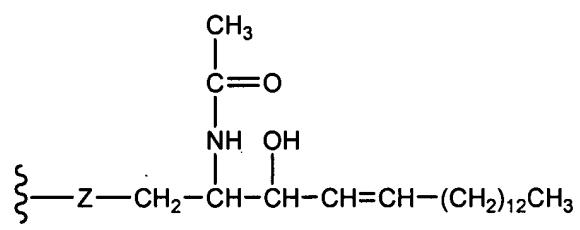
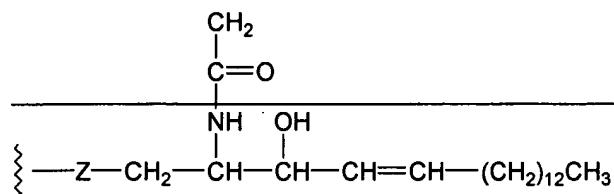
74. (Original) A compound of claim 62, wherein R⁴ is selected from the group consisting of:



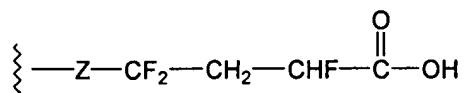
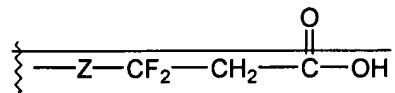
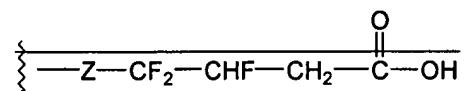
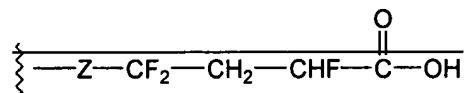
75. (Original) A compound of claim 62, wherein R⁴ is:

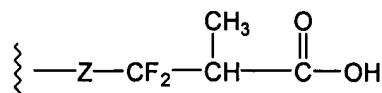
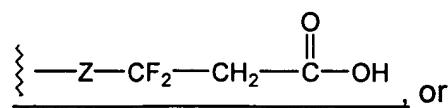
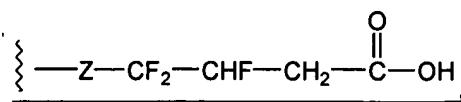


76. (Currently Amended) A compound of claim 62, wherein R⁴ is:

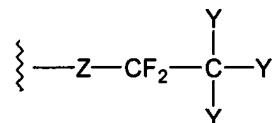


77. (Currently Amended) A compound of claim 62, wherein R⁴ is:

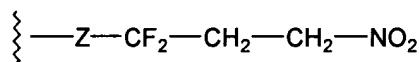




78. (Original) A compound of claim 62, wherein R⁴ is:



79. (Original) A compound of claim 62, wherein R⁴ is:



80. (Canceled).

81. (Canceled).

82. (Canceled).

83. (Canceled).

84. (Canceled).

85. (Canceled).

86. (Canceled).

87. (Canceled).

88. (Currently Amended) The method of claims 56 or 57 ~~86 or 87~~,
wherein the hyperproliferative cell is a cancer cell.

89. (Original) The method of claim 88, wherein the cancer cell is selected
from the group consisting of a colorectal cell, a head and neck cancer cell, a breast
cancer cell, a liver cancer cell and a gastric cancer cell.